



96TH GENERAL ASSEMBLY

State of Illinois

2009 and 2010

SB1506

Introduced 2/18/2009, by Sen. Jacqueline Y. Collins

SYNOPSIS AS INTRODUCED:

New Act
215 ILCS 5/155.36
215 ILCS 134/40
215 ILCS 134/45

Creates the Health Carrier External Review Act. Sets forth standards for independent external review procedures for adverse determinations by a health carrier against a covered person. Provides that the Act applies to an entity subject to the insurance laws and regulations of this State or subject to the jurisdiction of the Director and that contracts or offers to contract concerning any costs of health care. Requires health carriers to notify covered persons and their health care providers in writing of the covered person's right to request an external review as provided by the Act. Sets forth notice requirements. Provides that a request for an external review shall not be made until the covered person has exhausted the health carrier's internal grievance process. Sets forth requirements for standard external reviews and expedited external reviews. Provides that an external review decision is binding on the health carrier and binding on the covered person except to the extent the covered person has other remedies available. Sets forth minimum qualifications for independent review organizations and provides that the Director shall approve independent review organizations eligible to be assigned to conduct external reviews. Provides that each health carrier shall maintain written records of external review requests for each calendar year and submit a report to the Director by March 1 of each year. Provides that the health carrier shall be solely responsible for paying the cost of external reviews. Sets forth disclosure requirements. Amends the Managed Care Reform and Patient Rights Act to provide that an enrollee may appeal adverse decisions in accordance with the Health Carrier External Review Act. Deletes a provision concerning external independent review. Makes other changes. Contains a severability clause. Effective January 1, 2010.

LRB096 10769 RPM 20965 b

1 AN ACT concerning insurance.

2 **Be it enacted by the People of the State of Illinois,**
3 **represented in the General Assembly:**

4 Section 1. Short title. This Act may be cited as the Health
5 Carrier External Review Act.

6 Section 5. Purpose and intent. The purpose of this Act is
7 to provide uniform standards for the establishment and
8 maintenance of external review procedures to assure that
9 covered persons have the opportunity for an independent review
10 of an adverse determination or final adverse determination, as
11 defined in this Act.

12 Section 10. Definitions. For the purposes of this Act:

13 "Adverse determination" means a determination by a health
14 carrier or its designee utilization review organization that an
15 admission, availability of care, continued stay, or other
16 health care service that is a covered benefit has been reviewed
17 and, based upon the information provided, does not meet the
18 health carrier's requirements for medical necessity,
19 appropriateness, health care setting, level of care, or
20 effectiveness, and the requested service or payment for the
21 service is therefore denied, reduced, or terminated.

22 "Authorized representative" means:

1 (i) a person to whom a covered person has given express
2 written consent to represent the covered person in an
3 external review;

4 (ii) a person authorized by law to provide substituted
5 consent for a covered person;

6 (iii) a family member of the covered person; or

7 (iv) the covered person's health care provider.

8 "Clinical review criteria" means the written screening
9 procedures, decision abstracts, clinical protocols, and
10 practice guidelines used by a health carrier to determine the
11 necessity and appropriateness of health care services.

12 "Director" means the Director of the Division of Insurance
13 within the Illinois Department of Financial and Professional
14 Regulation.

15 "Covered benefits" or "benefits" means those health care
16 services to which a covered person is entitled under the terms
17 of a health benefit plan.

18 "Covered person" means a policyholder, subscriber,
19 enrollee, or other individual participating in a health benefit
20 plan.

21 "Emergency medical condition" means the sudden onset of a
22 health condition or illness that requires immediate medical
23 attention, where failure to provide medical attention would
24 result in a serious impairment to bodily functions, serious
25 dysfunction of a bodily organ or part, or would place the
26 person's health in serious jeopardy.

1 "Emergency services" means health care items and services
2 furnished or required to evaluate and treat an emergency
3 medical condition.

4 "Evidence-based standard" means a standard of care
5 developed through the judicious use of the current best
6 evidence and based on an overall systematic review of
7 applicable research.

8 "Facility" means an institution providing health care
9 services or a health care setting.

10 "Final adverse determination" means an adverse
11 determination involving a covered benefit that has been upheld
12 by a health carrier, or its designee utilization review
13 organization, at the completion of the health carrier's
14 internal grievance process procedures as set forth in Section
15 45 of the Managed Care Reform and Patient Rights Act.

16 "Health benefit plan" means a policy, contract,
17 certificate, plan, or agreement offered or issued by a health
18 carrier to provide, deliver, arrange for, pay for, or reimburse
19 any of the costs of health care services.

20 "Health care provider" or "provider" means a physician or
21 other health care practitioner licensed, accredited, or
22 certified to perform specified health care services consistent
23 with State law, responsible for recommending health care
24 services on behalf of a covered person.

25 "Health care services" means services for the diagnosis,
26 prevention, treatment, cure, or relief of a health condition,

1 illness, injury, or disease.

2 "Health carrier" means an entity subject to the insurance
3 laws and regulations of this State, or subject to the
4 jurisdiction of the Director, that contracts or offers to
5 contract to provide, deliver, arrange for, pay for, or
6 reimburse any of the costs of health care services, including a
7 sickness and accident insurance company, a health maintenance
8 organization, a nonprofit hospital and health service
9 corporation, or any other entity providing a plan of health
10 insurance, health benefits, or health care services. "Health
11 carrier" also means Limited Health Service Organizations
12 (LHSO) and Voluntary Health Service Plans.

13 "Health information" means information or data, whether
14 oral or recorded in any form or medium, and personal facts or
15 information about events or relationships that relate to:

16 (1) the past, present, or future physical, mental, or
17 behavioral health or condition of an individual or a member
18 of the individual's family;

19 (2) the provision of health care services to an
20 individual; or

21 (3) payment for the provision of health care services
22 to an individual.

23 "Independent review organization" means an entity that
24 conducts independent external reviews of adverse
25 determinations and final adverse determinations.

26 "Medical or scientific evidence" means evidence found in

1 the following sources:

2 (1) peer-reviewed scientific studies published in or
3 accepted for publication by medical journals that meet
4 nationally recognized requirements for scientific
5 manuscripts and that submit most of their published
6 articles for review by experts who are not part of the
7 editorial staff;

8 (2) peer-reviewed medical literature, including
9 literature relating to therapies reviewed and approved by a
10 qualified institutional review board, biomedical
11 compendia, and other medical literature that meet the
12 criteria of the National Institutes of Health's Library of
13 Medicine for indexing in Index Medicus (Medline) and
14 Elsevier Science Ltd. for indexing in Excerpta Medicus
15 (EMBASE);

16 (3) medical journals recognized by the Secretary of
17 Health and Human Services under Section 1861(t)(2) of the
18 federal Social Security Act;

19 (4) the following standard reference compendia:

20 (a) The American Hospital Formulary Service-Drug
21 Information;

22 (b) Drug Facts and Comparisons;

23 (c) The American Dental Association Accepted
24 Dental Therapeutics; and

25 (d) The United States Pharmacopoeia-Drug
26 Information;

1 (5) findings, studies, or research conducted by or
2 under the auspices of federal government agencies and
3 nationally recognized federal research institutes,
4 including:

5 (a) the federal Agency for Healthcare Research and
6 Quality;

7 (b) the National Institutes of Health;

8 (c) the National Cancer Institute;

9 (d) the National Academy of Sciences;

10 (e) the Centers for Medicare & Medicaid Services;

11 (f) the federal Food and Drug Administration; and

12 (g) any national board recognized by the National
13 Institutes of Health for the purpose of evaluating the
14 medical value of health care services; or

15 (6) any other medical or scientific evidence that is
16 comparable to the sources listed in items (1) through (5).

17 "Protected health information" means health information
18 (i) that identifies an individual who is the subject of the
19 information; or (ii) with respect to which there is a
20 reasonable basis to believe that the information could be used
21 to identify an individual.

22 "Utilization review" has the meaning provided by the
23 Managed Care Reform and Patient Rights Act.

24 "Utilization review organization" means a utilization
25 review program as defined by the Managed Care Reform and
26 Patient Rights Act.

1 Section 15. Applicability and scope.

2 (a) Except as provided in subsection (b), this Act shall
3 apply to all health carriers.

4 (b) The provisions of this Act shall not apply to a policy
5 or certificate that provides coverage only for a specified
6 disease, specified accident or accident-only coverage, credit,
7 dental, disability income, hospital indemnity, long-term care
8 insurance, as defined by Article XIXA of the Illinois Insurance
9 Code, vision care, or any other limited supplemental benefit or
10 to a Medicare supplement policy of insurance, as defined by the
11 Director by regulation, coverage under a plan through Medicare,
12 Medicaid, or the federal employees health benefits program, any
13 coverage issued under Chapter 55 of Title 10, U.S. Code and any
14 coverage issued as supplement to that coverage, any coverage
15 issued as supplemental to liability insurance, workers'
16 compensation or similar insurance, automobile medical-payment
17 insurance, or any insurance under which benefits are payable
18 with or without regard to fault, whether written on a group
19 blanket or individual basis.

20 Section 20. Notice of right to external review.

21 (a) At the same time the health carrier sends written
22 notice of a covered person's right to appeal a coverage
23 decision as provided by the Managed Care Reform and Patient
24 Rights Act, a health carrier shall notify a covered person and

1 a covered person's health care provider in writing of the
2 covered person's right to request an external review as
3 provided by this Act.

4 (1) The written notice required shall include the
5 following, or substantially equivalent, language: "We have
6 denied your request for the provision of or payment for a
7 health care service or course of treatment. You have the
8 right to have our decision reviewed by an independent
9 review organization not associated with us if our decision
10 involved making a judgment as to the medical necessity,
11 appropriateness, health care setting, level of care, or
12 effectiveness of the health care service or treatment you
13 requested by submitting a written request for an external
14 review to us. Upon receipt of your request an independent
15 review organization registered with the Department of
16 Financial and Professional Regulation, Division of
17 Insurance will be assigned to review our decision."

18 (2) The notice shall also include the appropriate
19 statements and information set forth in subsection (b) of
20 this Section.

21 (b) Expedited review prior to a final adverse
22 determination. The health carrier shall include in the notice
23 required under subsection (a) of this Section for a notice
24 related to an adverse determination, a statement informing the
25 covered person that:

26 (1) If the covered person has a medical condition where

1 the timeframe for completion of an expedited internal
2 review of a grievance involving an adverse determination
3 set forth in the Managed Care Reform and Patient Rights Act
4 would seriously jeopardize the life or health of the
5 covered person or would jeopardize the covered person's
6 ability to regain maximum function, the covered person or
7 the covered person's authorized representative may file a
8 request for an expedited external review.

9 (2) The covered person or the covered person's
10 authorized representative may file a request for an
11 expedited external review at the same time the covered
12 person or the covered person's authorized representative
13 files a request for an expedited internal appeal involving
14 an adverse determination as set forth in the Managed Care
15 Reform and Patient Rights Act, if the adverse determination
16 involves a denial of coverage based on a determination that
17 the recommended or requested health care service or
18 treatment is experimental or investigational and the
19 covered person's health care provider certifies in writing
20 that the recommended or requested health care service or
21 treatment that is the subject of the adverse determination
22 would be significantly less effective if not promptly
23 initiated. The independent review organization assigned to
24 conduct the expedited external review will determine
25 whether the covered person shall be required to complete
26 the expedited review of the grievance prior to conducting

1 the expedited external review.

2 (c) Expedited review upon final adverse determination. The
3 health carrier shall include in the notice required under
4 subsection (a) for a notice related to an adverse
5 determination, a statement informing the covered person that:

6 (1) if the covered person has a medical condition where
7 the timeframe for completion of a standard external review
8 would seriously jeopardize the life or health of the
9 covered person or would jeopardize the covered person's
10 ability to regain maximum function, then the covered person
11 or the covered person's authorized representative may file
12 a request for an expedited external review;

13 (2) if a final adverse determination concerns an
14 admission, availability of care, continued stay, or health
15 care service for which the covered person received
16 emergency services, but has not been discharged from a
17 facility, then the covered person, or the covered person's
18 authorized representative, may request an expedited
19 external review; or

20 (3) if a final adverse determination concerns a denial
21 of coverage based on a determination that the recommended
22 or requested health care service or treatment is
23 experimental or investigational, and the covered person's
24 health care provider certifies in writing that the
25 recommended or requested health care service or treatment
26 that is the subject of the request would be significantly

1 less effective if not promptly initiated, then the covered
2 person or the covered person's authorized representative
3 may request an expedited external review.

4 (d) In addition to the information to be provided pursuant
5 to subsections (a), (b), and (c) of this Section, the health
6 carrier shall include a copy of the description of both the
7 required standard and expedited external review procedures.
8 The description shall highlight the external review procedures
9 that give the covered person or the covered person's authorized
10 representative the opportunity to submit additional
11 information, including any forms used to process an external
12 review.

13 (e) In addition to the information to be provided under
14 subsections (a), (b), or (c) of this Section, the health
15 carrier shall include an authorization form that complies with
16 the requirements of the federal Health Insurance Portability
17 and Accountability Act (HIPAA) (45 CFR Section 164.508), by
18 which the covered person, for purposes of conducting an
19 external review under this Act, authorizes the health carrier
20 and the covered person's health care provider to disclose
21 protected health information, including medical records,
22 concerning the covered person that are pertinent to the
23 external review.

24 Section 25. Request for external review. A covered person
25 or the covered person's authorized representative may make a

1 request for an external or expedited external review of an
2 adverse determination or final adverse determination. Requests
3 under this Section shall be made directly to the health carrier
4 that made the adverse or final adverse determination. All
5 requests for external review shall be in writing except for
6 requests for expedited external reviews which may be made
7 orally. Health carriers must provide covered persons with forms
8 to request external reviews.

9 Section 30. Exhaustion of internal grievance process.
10 Except as provided in subsection (b) of Section 20 of this Act,
11 a request for an external review shall not be made until the
12 covered person has exhausted the health carrier's internal
13 grievance process as set forth in the Managed Care Reform and
14 Patient Rights Act. A covered person shall also be considered
15 to have exhausted the health carrier's internal grievance
16 process for purposes of this Section:

17 (a) If the covered person or the covered person's
18 authorized representative filed a request for an internal
19 review of an adverse determination pursuant to the Managed
20 Care Reform and Patient Rights Act and has not received a
21 written decision on the request from the health carrier
22 within 15 days, except to the extent the covered person or
23 the covered person's authorized representative requested
24 or agreed to a delay; or

25 (b) If the covered person or the covered person's

1 authorized representative filed a request for an expedited
2 internal review of an adverse determination pursuant to the
3 Managed Care Reform and Patient Rights Act and has not
4 received a decision on request from the health carrier
5 within 48 hours, except to the extent the covered person or
6 the covered person's authorized representative requested
7 or agreed to a delay.

8 A covered person need not exhaust a health carrier's
9 internal grievance procedures as set forth in the Managed Care
10 Reform and Patient Rights Act if the health carrier agrees to
11 waive the exhaustion requirement.

12 Section 35. Standard external review.

13 (a) Within 4 months after the date of receipt of a notice
14 of an adverse determination or final adverse determination, a
15 covered person or the covered person's authorized
16 representative may file a request for an external review with
17 the health carrier.

18 (b) Within 5 business days following the date of receipt of
19 the external review request, the health carrier shall complete
20 a preliminary review of the request to determine whether:

21 (1) the individual is or was a covered person in the
22 health benefit plan at the time the health care service was
23 requested or at the time the health care service was
24 provided;

25 (2) the health care service that is the subject of the

1 adverse determination or the final adverse determination
2 is a covered service under the covered person's health
3 benefit plan, but the health carrier has determined that
4 the health care service is not covered because it does not
5 meet the health carrier's requirements for medical
6 necessity, appropriateness, health care setting, level of
7 care, or effectiveness;

8 (3) the covered person has exhausted the health
9 carrier's internal grievance process as set forth in
10 Section 30 of this Act;

11 (4) for appeals relating to a determination based on
12 treatment being experimental or investigational, the
13 covered person's health care provider has certified that
14 one of the following situations is applicable:

15 (A) standard health care services or treatments
16 have not been effective in improving the condition of
17 the covered person;

18 (B) standard health care services or treatments
19 are not medically appropriate for the covered person;

20 (C) there is no available standard health care
21 service or treatment covered by the health carrier that
22 is more beneficial than the recommended or requested
23 health care service or treatment;

24 (D) the health care service or treatment is likely
25 to be more beneficial to the covered person, in the
26 health care provider's opinion, than any available

1 standard health care services or treatments; or

2 (E) that scientifically valid studies using
3 accepted protocols demonstrate that the health care
4 service or treatment requested is likely to be more
5 beneficial to the covered person than any available
6 standard health care services or treatments; and

7 (5) The covered person has attempted to provide all the
8 information and forms minimally required to process an
9 external review as specified in this Act.

10 (c) Within one business day after completion of the
11 preliminary review, the health carrier shall notify the covered
12 person, the covered person's health care provider, and, if
13 applicable, the covered person's authorized representative in
14 writing whether the request is complete and eligible for
15 external review. If the request:

16 (i) is not complete, the health carrier shall
17 inform the covered person, the covered person's health
18 care provider and, if applicable, the covered person's
19 authorized representative in writing and include in
20 the notice what information or materials are required
21 by this Act to make the request complete; or

22 (ii) is not eligible for external review, the
23 health carrier shall inform the covered person, the
24 covered person's health care provider, and if
25 applicable, the covered person's authorized
26 representative in writing and include in the notice the

1 reasons for its ineligibility.

2 The notice of initial determination of ineligibility
3 shall include a statement informing the covered person, the
4 covered person's health care provider and, if applicable,
5 the covered person's authorized representative that a
6 health carrier's initial determination that the external
7 review request is ineligible for review may be appealed to
8 the Director by filing a complaint with the Director.

9 Notwithstanding a health carrier's initial
10 determination that the request is ineligible and require
11 that it be referred for external review, the Director may
12 determine that a request is eligible for external review.

13 (d) Whenever a request is eligible for external review the
14 health carrier shall, within 3 business days:

15 (1) assign an independent review organization from the
16 list of approved independent review organizations compiled
17 and maintained by the Director; and

18 (2) notify in writing the covered person, the covered
19 person's health care provider and, if applicable, the
20 covered person's authorized representative of the
21 request's eligibility and acceptance for external review
22 and the name of the independent review organization.

23 The health carrier shall include in the notice provided to
24 the covered person, the covered person's health care provider
25 and, if applicable, the covered person's authorized
26 representative a statement that the covered person or the

1 covered person's authorized representative may, within 5
2 business days following the date of receipt of the notice
3 provided pursuant to item (1) of this subsection (d), submit in
4 writing to the assigned independent review organization
5 additional information that the independent review
6 organization shall consider when conducting the external
7 review. The independent review organization is not required to,
8 but may, accept and consider additional information submitted
9 after 5 business days.

10 (e) The assignment of an approved independent review
11 organization to conduct an external review in accordance with
12 this Section shall be done on a random basis among those
13 approved independent review organizations qualified to conduct
14 external review except for instances of conflict of interest
15 concerns pursuant to this Act.

16 (f) Upon assignment of an independent review organization,
17 the health carrier or its designee utilization review
18 organization shall, within 5 business days, provide to the
19 assigned independent review organization the documents and any
20 information considered in making the adverse determination or
21 final adverse determination.

22 (1) Except as provided in item (2) of this subsection
23 (f), failure by the health carrier or its utilization
24 review organization to provide the documents and
25 information within the specified time frame shall not delay
26 the conduct of the external review.

1 (2) If the health carrier or its utilization review
2 organization fails to provide the documents and
3 information within the specified time frame, the assigned
4 independent review organization may terminate the external
5 review and make a decision to reverse the adverse
6 determination or final adverse determination.

7 (3) Within one business day after making the decision
8 to terminate the external review and make a decision to
9 reverse the adverse determination or final adverse
10 determination under item (2) of this subsection (f), the
11 independent review organization shall notify the health
12 carrier, the covered person, the covered person's health
13 care provider and, if applicable, the covered person's
14 authorized representative, of its decision to reverse the
15 adverse determination.

16 (g) Upon receipt of the information from the health carrier
17 or its utilization review organization, the assigned
18 independent review organization shall review all of the
19 information and documents and any other information submitted
20 in writing to the independent review organization by the
21 covered person and the covered person's authorized
22 representative.

23 (h) Upon receipt of any information submitted by the
24 covered person or the covered person's authorized
25 representative, the independent review organization shall
26 forward the information to the health carrier within 1 business

1 day.

2 (1) Upon receipt of the information, if any, the health
3 carrier may reconsider its adverse determination or final
4 adverse determination that is the subject of the external
5 review.

6 (2) Reconsideration by the health carrier of its
7 adverse determination or final adverse determination shall
8 not delay or terminate the external review.

9 (3) The external review may only be terminated if the
10 health carrier decides, upon completion of its
11 reconsideration, to reverse its adverse determination or
12 final adverse determination and provide coverage or
13 payment for the health care service that is the subject of
14 the adverse determination or final adverse determination.
15 In such cases, the following provisions shall apply:

16 (A) Within one business day after making the
17 decision to reverse its adverse determination or final
18 adverse determination, the health carrier shall notify
19 the covered person, the covered person's health care
20 provider, if applicable, the covered person's
21 authorized representative, and the assigned
22 independent review organization in writing of its
23 decision.

24 (B) Upon notice from the health carrier that the
25 health carrier has made a decision to reverse its
26 adverse determination or final adverse determination,

1 the assigned independent review organization shall
2 terminate the external review.

3 (i) In addition to the documents and information provided
4 by the health carrier or its utilization review organization
5 and the covered person and the covered person's authorized
6 representative, if any, the independent review organization,
7 to the extent the information or documents are available and
8 the independent review organization considers them
9 appropriate, shall consider the following in reaching a
10 decision:

11 (1) the covered person's pertinent medical records;

12 (2) the covered person's health care provider's
13 recommendation;

14 (3) consulting reports from appropriate health care
15 providers and other documents submitted by the health
16 carrier, the covered person, and the covered person's
17 authorized representative;

18 (4) the terms of coverage under the covered person's
19 health benefit plan with the health carrier to ensure that
20 the health care service or treatment that is the subject of
21 the opinion is experimental or investigational would
22 otherwise be covered under the terms of coverage of the
23 covered person's health benefit plan with the health
24 carrier;

25 (5) the most appropriate practice guidelines, which
26 shall include applicable evidence-based standards and may

1 include any other practice guidelines developed by the
2 federal government, national or professional medical
3 societies, boards, and associations;

4 (6) any applicable clinical review criteria developed
5 and used by the health carrier or its designee utilization
6 review organization; and

7 (7) the opinion of the independent review
8 organization's clinical reviewer or reviewers after
9 considering items (1) through (6) of this subsection (i) to
10 the extent the information or documents are available and
11 the clinical reviewer or reviewers considers the
12 information or documents relevant.

13 (j) Within 5 days after the date of receipt of all
14 necessary information, the assigned independent review
15 organization shall provide written notice of its decision to
16 uphold or reverse the adverse determination or the final
17 adverse determination to the health carrier, the covered
18 person, the covered person's health care provider and, if
19 applicable, the covered person's authorized representative. In
20 such cases, the following provisions shall apply:

21 (1) The independent review organization shall include
22 in the notice:

23 (A) a general description of the reason for the
24 request for external review;

25 (B) the date the independent review organization
26 received the assignment from the health carrier to

1 conduct the external review;

2 (C) the time period during which the external
3 review was conducted;

4 (D) references to the evidence or documentation,
5 including the evidence-based standards, considered in
6 reaching its decision.

7 (E) the date of its decision; and

8 (F) the principal reason or reasons for its
9 decision, including what applicable, if any,
10 evidence-based standards that were a basis for its
11 decision.

12 (2) For reviews of experimental or investigational
13 treatments, the notice shall include the following
14 information:

15 (A) a description of the covered person's medical
16 condition;

17 (B) a description of the indicators relevant to
18 whether there is sufficient evidence to demonstrate
19 that the recommended or requested health care service
20 or treatment is more likely than not to be more
21 beneficial to the covered person than any available
22 standard health care services or treatments and the
23 adverse risks of the recommended or requested health
24 care service or treatment would not be substantially
25 increased over those of available standard health care
26 services or treatments;

1 (C) a description and analysis of any medical or
2 scientific evidence considered in reaching the
3 opinion;

4 (D) a description and analysis of any
5 evidence-based standards; and

6 (E) whether the recommended or requested health
7 care service or treatment has been approved by the
8 federal Food and Drug Administration, for the
9 condition; or

10 (F) Whether medical or scientific evidence or
11 evidence-based standards demonstrate that the expected
12 benefits of the recommended or requested health care
13 service or treatment is more likely than not to be more
14 beneficial to the covered person than any available
15 standard health care service or treatment and the
16 adverse risks of the recommended or requested health
17 care service or treatment would not be substantially
18 increased over those of available standard health care
19 services or treatments; in reaching a decision, the
20 assigned independent review organization is not bound
21 by any decisions or conclusions reached during the
22 health carrier's utilization review process or the
23 health carrier's internal grievance or appeals
24 process.

25 (3) Upon receipt of a notice of a decision reversing
26 the adverse determination or final adverse determination,

1 the health carrier immediately shall approve the coverage
2 that was the subject of the adverse determination or final
3 adverse determination.

4 Section 40. Expedited external review.

5 (a) A covered person or a covered person's authorized
6 representative may file a request for an expedited external
7 review with the health carrier either orally or in writing;

8 (1) immediately after the date of receipt of a notice a
9 final adverse determination as provided by subsection (c)
10 of Section 20; or

11 (2) if a health carrier fails to provide a decision on
12 request for an expedited internal appeal within 48 hours as
13 provided by subsection (b) of Section 30.

14 (b) Upon receipt of a request for an expedited external
15 review as provided in subsections (b) and (c) of Section 20,
16 the health carrier shall immediately assign an independent
17 review organization from the list of approved independent
18 review organizations compiled and maintained by the Director to
19 conduct the expedited review. In such cases, the following
20 provisions shall apply:

21 (1) The assignment by the health carrier of an approved
22 independent review organization to conduct an external
23 review in accordance with this Section shall be done on a
24 random basis among those approved independent review
25 organizations except as may be prohibited by conflict of

1 interest concerns pursuant to Section 60 of this Act.

2 (2) Immediately upon assigning an independent review
3 organization to perform an expedited external review, but
4 in no case less than 24 hours after assigning the
5 independent review organization, the health carrier or its
6 designee utilization review organization shall provide or
7 transmit all necessary documents and information
8 considered in making the final adverse determination to the
9 assigned independent review organization electronically or
10 by telephone or facsimile or any other available
11 expeditious method.

12 (3) If the health carrier or its utilization review
13 organization fails to provide the documents and
14 information within the specified time frame, the assigned
15 independent review organization may terminate the external
16 review and make a decision to reverse the adverse
17 determination or final adverse determination.

18 (4) Within one business day after making the decision
19 to terminate the external review and make a decision to
20 reverse the adverse determination or final adverse
21 determination under item (2) of this subsection (b), the
22 independent review organization shall notify the health
23 carrier, the covered person, the covered person's health
24 care provider and, if applicable, the covered person's
25 authorized representative of its decision to reverse the
26 adverse determination.

1 (c) In addition to the documents and information provided
2 by the health carrier or its utilization review organization
3 and any documents and information provided by the covered
4 person and the covered person's authorized representative, the
5 independent review organization shall consider the following
6 in reaching a decision:

7 (1) the covered person's pertinent medical records;

8 (2) the covered person's health care provider's
9 recommendation;

10 (3) consulting reports from appropriate health care
11 providers and other documents submitted by the health
12 carrier, the covered person and the covered person's
13 authorized representative;

14 (4) the terms of coverage under the covered person's
15 health benefit plan with the health carrier to ensure that
16 the health care service or treatment that is the subject of
17 the opinion is experimental or investigational would
18 otherwise be covered under the terms of coverage of the
19 covered person's health benefit plan with the health
20 carrier;

21 (5) the most appropriate practice guidelines, which
22 shall include applicable evidence-based standards and may
23 include any other practice guidelines developed by the
24 federal government, national or professional medical
25 societies, boards, and associations;

26 (6) any applicable clinical review criteria developed

1 and used by the health carrier or its designee utilization
2 review organization; and

3 (7) whether for experimental or investigational
4 denials:

5 (A) the recommended or requested health care
6 service or treatment has been approved by the federal
7 Food and Drug Administration, if applicable, for the
8 condition; or

9 (B) medical or scientific evidence or
10 evidence-based standards demonstrate that the expected
11 benefits of the recommended or requested health care
12 service or treatment is more likely than not to be
13 beneficial to the covered person than any available
14 standard health care service or treatment and the
15 adverse risks of the recommended or requested health
16 care service or treatment would not be substantially
17 increased over those of available standard health care
18 services or treatments.

19 (d) As expeditiously as the covered person's medical
20 condition or circumstances requires, but in no event more than
21 48 hours after the receipt of all pertinent information, the
22 assigned independent review organization shall:

23 (1) make a decision to uphold or reverse the final
24 adverse determination; and

25 (2) notify the health carrier, the covered person, the
26 covered person's health care provider, and if applicable,

1 the covered person's authorized representative, of the
2 decision.

3 In reaching a decision, the assigned independent review
4 organization is not bound by any decisions or conclusions
5 reached during the health carrier's utilization review process
6 or the health carrier's internal grievance process as set forth
7 in the Managed Care Reform and Patient Rights Act.

8 Upon receipt of notice of a decision reversing the final
9 adverse determination, the health carrier shall immediately
10 approve the coverage that was the subject of the final adverse
11 determination. Within 48 hours after the date of providing the
12 notice required in this subsection (d), the assigned
13 independent review organization shall provide written
14 confirmation of the decision to the health carrier, the covered
15 person, the covered person's health care provider, and if
16 applicable, the covered person's authorized representative
17 including:

18 (A) a general description of the reason for the
19 request for external review;

20 (B) the date the independent review organization
21 received the assignment from the health carrier to
22 conduct the external review;

23 (C) the date the external review was conducted;

24 (D) the date of its decision;

25 (E) the principal reason or reasons for its
26 decision, including what applicable, if any,

1 evidence-based standards were a basis for its
2 decision; and

3 (F) references to the evidence or documentation,
4 including the evidence-based standards, considered in
5 reaching its decision.

6 Section 45. Binding nature of external review decision. An
7 external review decision is binding on the health carrier. An
8 external review decision is binding on the covered person
9 except to the extent the covered person has other remedies
10 available under applicable federal or State law. A covered
11 person or the covered person's authorized representative may
12 not file a subsequent request for external review involving the
13 same adverse determination or final adverse determination for
14 which the covered person has already received an external
15 review decision pursuant to this Act.

16 Section 50. Approval of independent review organizations.

17 (a) The Director shall approve independent review
18 organizations eligible to be assigned to conduct external
19 reviews under this Act.

20 (b) In order to be eligible for approval by the Director
21 under this Section to conduct external reviews under this Act
22 an independent review organization:

23 (1) except as otherwise provided in this Section, shall
24 be accredited by a nationally recognized private

1 accrediting entity that the Director has determined has
2 independent review organization accreditation standards
3 that are equivalent to or exceed the minimum qualifications
4 for independent review; and

5 (2) shall submit an application for approval in
6 accordance with subsection (d) of this Section.

7 (c) The Director shall develop an application form for
8 initially approving and for reapproving independent review
9 organizations to conduct external reviews.

10 (d) Any independent review organization wishing to be
11 approved to conduct external reviews under this Act shall
12 submit the application form and include with the form all
13 documentation and information necessary for the Director to
14 determine if the independent review organization satisfies the
15 minimum qualifications established under this Act. The
16 Director may:

17 (1) approve independent review organizations that are
18 not accredited by a nationally recognized private
19 accrediting entity if there are no acceptable nationally
20 recognized private accrediting entities providing
21 independent review organization accreditation; and

22 (2) by rule establish an application fee that
23 independent review organizations shall submit to the
24 Director with an application for approval and renewing.

25 (e) An approval is effective for 2 years, unless the
26 Director determines before its expiration that the independent

1 review organization is not satisfying the minimum
2 qualifications established under this Act.

3 (f) Whenever the Director determines that an independent
4 review organization has lost its accreditation or no longer
5 satisfies the minimum requirements established under this Act,
6 the Director shall terminate the approval of the independent
7 review organization and remove the independent review
8 organization from the list of independent review organizations
9 approved to conduct external reviews under this Act that is
10 maintained by the Director.

11 (g) The Director shall maintain and periodically update a
12 list of approved independent review organizations.

13 (h) The Director may promulgate regulations to carry out
14 the provisions of this Section.

15 Section 55. Minimum qualifications for independent Review
16 organizations.

17 (a) To be approved to conduct external reviews, an
18 independent review organization shall have and maintain
19 written policies and procedures that govern all aspects of both
20 the standard external review process and the expedited external
21 review process set forth in this Act that include, at a
22 minimum:

23 (1) a quality assurance mechanism that ensures that:

24 (A) external reviews are conducted within the
25 specified time frames and required notices are

1 provided in a timely manner;

2 (B) selection of qualified and impartial clinical
3 reviewers to conduct external reviews on behalf of the
4 independent review organization and suitable matching
5 of reviewers to specific cases and that the independent
6 review organization employs or contracts with an
7 adequate number of clinical reviewers to meet this
8 objective;

9 (C) in assigning clinical reviewers, the
10 independent review organization selects physicians or
11 other health care professionals who, through clinical
12 experience in the past 3 years, are experts in the
13 treatment of the covered person's condition and
14 knowledgeable about the recommended or requested
15 health care service or treatment;

16 (D) the health carrier, the covered person and the
17 covered person's authorized representative shall not
18 choose or control the choice of the physicians or other
19 health care professionals to be selected to conduct the
20 external review;

21 (E) confidentiality of medical and treatment
22 records and clinical review criteria; and

23 (F) any person employed by or under contract with
24 the independent review organization adheres to the
25 requirements of this Act;

26 (2) a toll-free telephone service operating on a

1 24-hour-day, 7-day-a-week basis that accepts, receives,
2 and records information related to external reviews and
3 provides appropriate instructions; and

4 (3) an agreement to maintain and provide to the
5 Director the information set out in Section 70 of this Act.

6 (b) All clinical reviewers assigned by an independent
7 review organization to conduct external reviews shall be
8 physicians or other appropriate health care providers who meet
9 the following minimum qualifications:

10 (1) be an expert in the treatment of the covered
11 person's medical condition that is the subject of the
12 external review;

13 (2) be knowledgeable about the recommended health care
14 service or treatment through recent or current actual
15 clinical experience treating patients with the same or
16 similar medical condition of the covered person;

17 (3) hold a non-restricted license in a state of the
18 United States and, for physicians, a current certification
19 by a recognized American medical specialty board in the
20 area or areas appropriate to the subject of the external
21 review; and

22 (4) have no history of disciplinary actions or
23 sanctions, including loss of staff privileges or
24 participation restrictions, that have been taken or are
25 pending by any hospital, governmental agency or unit, or
26 regulatory body that raise a substantial question as to the

1 clinical reviewer's physical, mental, or professional
2 competence or moral character.

3 (c) In addition to the requirements set forth in subsection
4 (a), an independent review organization may not own or control,
5 be a subsidiary of, or in any way be owned, or controlled by,
6 or exercise control with a health benefit plan, a national,
7 State, or local trade association of health benefit plans, or a
8 national, State, or local trade association of health care
9 providers.

10 (d) Conflicts of interest prohibited. In addition to the
11 requirements set forth in subsections (a), (b), and (c) of this
12 Section, to be approved pursuant to this Act to conduct an
13 external review of a specified case, neither the independent
14 review organization selected to conduct the external review nor
15 any clinical reviewer assigned by the independent organization
16 to conduct the external review may have a material
17 professional, familial or financial conflict of interest with
18 any of the following:

19 (1) the health carrier that is the subject of the
20 external review;

21 (2) the covered person whose treatment is the subject
22 of the external review or the covered person's authorized
23 representative;

24 (3) any officer, director or management employee of the
25 health carrier that is the subject of the external review;

26 (4) the health care provider, the health care

1 provider's medical group or independent practice
2 association recommending the health care service or
3 treatment that is the subject of the external review;

4 (5) the facility at which the recommended health care
5 service or treatment would be provided; or

6 (6) The developer or manufacturer of the principal
7 drug, device, procedure, or other therapy being
8 recommended for the covered person whose treatment is the
9 subject of the external review.

10 (e) An independent review organization that is accredited
11 by a nationally recognized private accrediting entity that has
12 independent review accreditation standards that the Director
13 has determined are equivalent to or exceed the minimum
14 qualifications of this Section shall be presumed to be in
15 compliance with this Section and shall be eligible for approval
16 under this Section.

17 (f) An independent review organization shall be unbiased.
18 An independent review organization shall establish and
19 maintain written procedures to ensure that it is unbiased in
20 addition to any other procedures required under this Section.

21 Section 60. Hold harmless for independent review
22 organizations. No independent review organization or clinical
23 reviewer working on behalf of an independent review
24 organization or an employee, agent or contractor of an
25 independent review organization shall be liable for damages to

1 any person for any opinions rendered or acts or omissions
2 performed within the scope of the organization's or person's
3 duties under the law during or upon completion of an external
4 review conducted pursuant to this Act, unless the opinion was
5 rendered or act or omission performed in bad faith or involved
6 gross negligence.

7 Section 65. External review reporting requirements.

8 (a) Each health carrier shall maintain written records in
9 the aggregate on all requests for external review for each
10 calendar year and submit a report to the Director in the format
11 specified by the Director by March 1 of each year.

12 (b) The report shall include in the aggregate:

13 (1) the total number of requests for external review;

14 (2) the total number of requests for expedited external
15 review;

16 (3) the total number of requests for external review
17 denied;

18 (4) the number of requests for external review
19 resolved, including:

20 (A) the number of requests for external review
21 resolved upholding the adverse determination or final
22 adverse determination;

23 (B) the number of requests for external review
24 resolved reversing the adverse determination or final
25 adverse determination;

1 (C) the number of requests for expedited external
2 review resolved upholding the adverse determination or
3 final adverse determination; and

4 (D) the number of requests for expedited external
5 review resolved reversing the adverse determination or
6 final adverse determination;

7 (5) the average length of time for resolution for an
8 external review;

9 (6) the average length of time for resolution for an
10 expedited external review;

11 (7) a summary of the types of coverages or cases for
12 which an external review was sought, as specified below:

13 (A) denial of care or treatment (dissatisfaction
14 regarding prospective non-authorization of a request
15 for care or treatment recommended by a provider
16 excluding diagnostic procedures and referral requests;
17 partial approvals and care terminations are also
18 considered to be denials);

19 (B) denial of diagnostic procedure
20 (dissatisfaction regarding prospective
21 non-authorization of a request for a diagnostic
22 procedure recommended by a provider; partial approvals
23 are also considered to be denials);

24 (C) denial of referral request (dissatisfaction
25 regarding non-authorization of a request for a
26 referral to another provider recommended by a PCP);

1 (D) claims and utilization review (dissatisfaction
2 regarding the concurrent or retrospective evaluation
3 of the coverage, medical necessity, efficiency or
4 appropriateness of health care services or treatment
5 plans; prospective "Denials of care or treatment,"
6 "Denials of diagnostic procedures" and "Denials of
7 referral requests" should not be classified in this
8 category, but the appropriate one above);

9 (8) the number of external reviews that were terminated
10 as the result of a reconsideration by the health carrier of
11 its adverse determination or final adverse determination
12 after the receipt of additional information from the
13 covered person or the covered person's authorized
14 representative; and

15 (9) any other information the Director may request or
16 require.

17 Section 70. Funding of external review. The health carrier
18 shall be solely responsible for paying the cost of external
19 reviews conducted by independent review organizations.

20 Section 75. Disclosure requirements.

21 (a) Each health carrier shall include a description of the
22 external review procedures in, or attached to, the policy,
23 certificate, membership booklet, and outline of coverage or
24 other evidence of coverage it provides to covered persons.

1 (b) The description required under subsection (a) of this
2 Section shall include a statement that informs the covered
3 person of the right of the covered person to file a request for
4 an external review of an adverse determination or final adverse
5 determination with the health carrier. The statement shall
6 explain that external review is available when the adverse
7 determination or final adverse determination involves an issue
8 of medical necessity, appropriateness, health care setting,
9 level of care, or effectiveness. The statement shall include
10 the toll-free telephone number and address of the Office of
11 Consumer Health Insurance within the Division of Insurance.

12 (c) In addition to subsection (b) of this Section, the
13 statement shall inform the covered person that, when filing a
14 request for an external review, the covered person will be
15 required to authorize the release of any medical records of the
16 covered person that may be required to be reviewed for the
17 purpose of reaching a decision on the external review.

18 Section 90. The Illinois Insurance Code is amended by
19 changing Sections 155.36 as follows:

20 (215 ILCS 5/155.36)

21 Sec. 155.36. Managed Care Reform and Patient Rights Act.
22 Insurance companies that transact the kinds of insurance
23 authorized under Class 1(b) or Class 2(a) of Section 4 of this
24 Code shall comply with Sections 45, ~~Section~~ 85 and the

1 definition of the term "emergency medical condition" in Section
2 10 of the Managed Care Reform and Patient Rights Act.

3 (Source: P.A. 91-617, eff. 1-1-00.)

4 Section 95. The Managed Care Reform and Patient Rights Act
5 is amended by changing Sections 40 and 45 as follows:

6 (215 ILCS 134/40)

7 Sec. 40. Access to specialists.

8 (a) All health care plans that require each enrollee to
9 select a health care provider for any purpose including
10 coordination of care shall permit an enrollee to choose any
11 available primary care physician licensed to practice medicine
12 in all its branches participating in the health care plan for
13 that purpose. The health care plan shall provide the enrollee
14 with a choice of licensed health care providers who are
15 accessible and qualified. Nothing in this Act shall be
16 construed to prohibit a health care plan from requiring a
17 health care provider to meet the health care plan's criteria in
18 order to coordinate access to health care.

19 (b) A health care plan shall establish a procedure by which
20 an enrollee who has a condition that requires ongoing care from
21 a specialist physician or other health care provider may apply
22 for a standing referral to a specialist physician or other
23 health care provider if a referral to a specialist physician or
24 other health care provider is required for coverage. The

1 application shall be made to the enrollee's primary care
2 physician. This procedure for a standing referral must specify
3 the necessary criteria and conditions that must be met in order
4 for an enrollee to obtain a standing referral. A standing
5 referral shall be effective for the period necessary to provide
6 the referred services or one year, except in the event of
7 termination of a contract or policy in which case Section 25 on
8 transition of services shall apply, if applicable. A primary
9 care physician may renew and re-renew a standing referral.

10 (c) The enrollee may be required by the health care plan to
11 select a specialist physician or other health care provider who
12 has a referral arrangement with the enrollee's primary care
13 physician or to select a new primary care physician who has a
14 referral arrangement with the specialist physician or other
15 health care provider chosen by the enrollee. If a health care
16 plan requires an enrollee to select a new physician under this
17 subsection, the health care plan must provide the enrollee with
18 both options provided in this subsection. When a participating
19 specialist with a referral arrangement is not available, the
20 primary care physician, in consultation with the enrollee,
21 shall arrange for the enrollee to have access to a qualified
22 participating health care provider, and the enrollee shall be
23 allowed to stay with his or her primary care physician. If a
24 secondary referral is necessary, the specialist physician or
25 other health care provider shall advise the primary care
26 physician. The primary care physician shall be responsible for

1 making the secondary referral. In addition, the health care
2 plan shall require the specialist physician or other health
3 care provider to provide regular updates to the enrollee's
4 primary care physician.

5 (d) When the type of specialist physician or other health
6 care provider needed to provide ongoing care for a specific
7 condition is not represented in the health care plan's provider
8 network, the primary care physician shall arrange for the
9 enrollee to have access to a qualified non-participating health
10 care provider within a reasonable distance and travel time at
11 no additional cost beyond what the enrollee would otherwise pay
12 for services received within the network. The referring
13 physician shall notify the plan when a referral is made outside
14 the network.

15 (e) The enrollee's primary care physician shall remain
16 responsible for coordinating the care of an enrollee who has
17 received a standing referral to a specialist physician or other
18 health care provider. If a secondary referral is necessary, the
19 specialist physician or other health care provider shall advise
20 the primary care physician. The primary care physician shall be
21 responsible for making the secondary referral. In addition, the
22 health care plan shall require the specialist physician or
23 other health care provider to provide regular updates to the
24 enrollee's primary care physician.

25 (f) If an enrollee's application for any referral is
26 denied, an enrollee may appeal the decision through the health

1 care plan's external independent review process in accordance
2 with as provided by the Illinois Health Carrier External Review
3 Act subsection (f) of Section 45 of this Act.

4 (g) Nothing in this Act shall be construed to require an
5 enrollee to select a new primary care physician when no
6 referral arrangement exists between the enrollee's primary
7 care physician and the specialist selected by the enrollee and
8 when the enrollee has a long-standing relationship with his or
9 her primary care physician.

10 (h) In promulgating rules to implement this Act, the
11 Department shall define "standing referral" and "ongoing
12 course of treatment".

13 (Source: P.A. 91-617, eff. 1-1-00.)

14 (215 ILCS 134/45)

15 Sec. 45. Health care services appeals and ~~7~~ complaints~~7~~ ~~and~~
16 ~~external independent reviews.~~

17 (a) A health care plan shall establish and maintain an
18 appeals procedure as outlined in this Act. Compliance with this
19 Act's appeals procedures shall satisfy a health care plan's
20 obligation to provide appeal procedures under any other State
21 law or rules. All appeals of a health care plan's
22 administrative determinations and complaints regarding its
23 administrative decisions shall be handled as required under
24 Section 50.

25 (b) When an appeal concerns a decision or action by a

1 health care plan, its employees, or its subcontractors that
2 relates to (i) health care services, including, but not limited
3 to, procedures or treatments, for an enrollee with an ongoing
4 course of treatment ordered by a health care provider, the
5 denial of which could significantly increase the risk to an
6 enrollee's health, or (ii) a treatment referral, service,
7 procedure, or other health care service, the denial of which
8 could significantly increase the risk to an enrollee's health,
9 the health care plan must allow for the filing of an appeal
10 either orally or in writing. Upon submission of the appeal, a
11 health care plan must notify the party filing the appeal, as
12 soon as possible, but in no event more than 24 hours after the
13 submission of the appeal, of all information that the plan
14 requires to evaluate the appeal. The health care plan shall
15 render a decision on the appeal within 24 hours after receipt
16 of the required information. The health care plan shall notify
17 the party filing the appeal and the enrollee, enrollee's
18 primary care physician, and any health care provider who
19 recommended the health care service involved in the appeal of
20 its decision orally followed-up by a written notice of the
21 determination.

22 (c) For all appeals related to health care services
23 including, but not limited to, procedures or treatments for an
24 enrollee and not covered by subsection (b) above, the health
25 care plan shall establish a procedure for the filing of such
26 appeals. Upon submission of an appeal under this subsection, a

1 health care plan must notify the party filing an appeal, within
2 3 business days, of all information that the plan requires to
3 evaluate the appeal. The health care plan shall render a
4 decision on the appeal within 15 business days after receipt of
5 the required information. The health care plan shall notify the
6 party filing the appeal, the enrollee, the enrollee's primary
7 care physician, and any health care provider who recommended
8 the health care service involved in the appeal orally of its
9 decision followed-up by a written notice of the determination.

10 (d) An appeal under subsection (b) or (c) may be filed by
11 the enrollee, the enrollee's designee or guardian, the
12 enrollee's primary care physician, or the enrollee's health
13 care provider. A health care plan shall designate a clinical
14 peer to review appeals, because these appeals pertain to
15 medical or clinical matters and such an appeal must be reviewed
16 by an appropriate health care professional. No one reviewing an
17 appeal may have had any involvement in the initial
18 determination that is the subject of the appeal. The written
19 notice of determination required under subsections (b) and (c)
20 shall include (i) clear and detailed reasons for the
21 determination, (ii) the medical or clinical criteria for the
22 determination, which shall be based upon sound clinical
23 evidence and reviewed on a periodic basis, and (iii) in the
24 case of an adverse determination, the procedures for requesting
25 an external independent review as provided by the Illinois
26 Health Carrier External Review Act ~~under subsection (f).~~

1 (e) If an appeal filed under subsection (b) or (c) is
2 denied for a reason including, but not limited to, the service,
3 procedure, or treatment is not viewed as medically necessary,
4 denial of specific tests or procedures, denial of referral to
5 specialist physicians or denial of hospitalization requests or
6 length of stay requests, any involved party may request an
7 external independent review as provided by the Illinois Health
8 Carrier External Review Act ~~under subsection (f) of the adverse~~
9 ~~determination.~~

10 ~~(f) External independent review.~~

11 ~~(1) The party seeking an external independent review~~
12 ~~shall so notify the health care plan. The health care plan~~
13 ~~shall seek to resolve all external independent reviews in~~
14 ~~the most expeditious manner and shall make a determination~~
15 ~~and provide notice of the determination no more than 24~~
16 ~~hours after the receipt of all necessary information when a~~
17 ~~delay would significantly increase the risk to an~~
18 ~~enrollee's health or when extended health care services for~~
19 ~~an enrollee undergoing a course of treatment prescribed by~~
20 ~~a health care provider are at issue.~~

21 ~~(2) Within 30 days after the enrollee receives written~~
22 ~~notice of an adverse determination, if the enrollee decides~~
23 ~~to initiate an external independent review, the enrollee~~
24 ~~shall send to the health care plan a written request for an~~
25 ~~external independent review, including any information or~~
26 ~~documentation to support the enrollee's request for the~~

1 ~~covered service or claim for a covered service.~~

2 ~~(3) Within 30 days after the health care plan receives~~
3 ~~a request for an external independent review from an~~
4 ~~enrollee, the health care plan shall:~~

5 ~~(A) provide a mechanism for joint selection of an~~
6 ~~external independent reviewer by the enrollee, the~~
7 ~~enrollee's physician or other health care provider,~~
8 ~~and the health care plan; and~~

9 ~~(B) forward to the independent reviewer all~~
10 ~~medical records and supporting documentation~~
11 ~~pertaining to the case, a summary description of the~~
12 ~~applicable issues including a statement of the health~~
13 ~~care plan's decision, the criteria used, and the~~
14 ~~medical and clinical reasons for that decision.~~

15 ~~(4) Within 5 days after receipt of all necessary~~
16 ~~information, the independent reviewer shall evaluate and~~
17 ~~analyze the case and render a decision that is based on~~
18 ~~whether or not the health care service or claim for the~~
19 ~~health care service is medically appropriate. The decision~~
20 ~~by the independent reviewer is final. If the external~~
21 ~~independent reviewer determines the health care service to~~
22 ~~be medically appropriate, the health care plan shall pay~~
23 ~~for the health care service.~~

24 ~~(5) The health care plan shall be solely responsible~~
25 ~~for paying the fees of the external independent reviewer~~
26 ~~who is selected to perform the review.~~

1 ~~(6) An external independent reviewer who acts in good~~
2 ~~faith shall have immunity from any civil or criminal~~
3 ~~liability or professional discipline as a result of acts or~~
4 ~~omissions with respect to any external independent review,~~
5 ~~unless the acts or omissions constitute wilful and wanton~~
6 ~~misconduct. For purposes of any proceeding, the good faith~~
7 ~~of the person participating shall be presumed.~~

8 ~~(7) Future contractual or employment action by the~~
9 ~~health care plan regarding the patient's physician or other~~
10 ~~health care provider shall not be based solely on the~~
11 ~~physician's or other health care provider's participation~~
12 ~~in this procedure.~~

13 ~~(8) For the purposes of this Section, an external~~
14 ~~independent reviewer shall:~~

15 ~~(A) be a clinical peer;~~

16 ~~(B) have no direct financial interest in~~
17 ~~connection with the case; and~~

18 ~~(C) have not been informed of the specific identity~~
19 ~~of the enrollee.~~

20 ~~(9) Nothing in this Section shall be construed to require a~~
21 ~~health care plan to pay for a health care service not covered~~
22 ~~under the enrollee's certificate of coverage or policy.~~

23 (Source: P.A. 91-617, eff. 1-1-00.)

24 Section 97. Severability. The provisions of this Act are
25 severable under Section 1.31 of the Statute on Statutes.

1 Section 99. Effective date. This Act takes effect January
2 1, 2010.